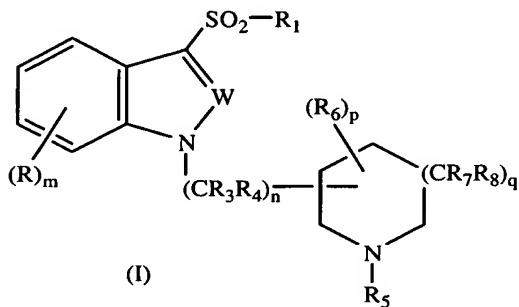


AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended) A compound of formula I



wherein

W is [[N or]] CR₂;

R is halogen, CN, OCO₂R₉, CO₂R₁₀, CONR₁₁R₁₂, SO_xR₁₃, NR₁₄R₁₅, OR₁₆, COR₁₇ or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₇cycloalkyl, aryl or heteroaryl group each optionally substituted;

R₁ is an optionally substituted C₁-C₆alkyl, C₃-C₇cycloalkyl, aryl, or heteroaryl group or an optionally substituted 8- to 13-membered bicyclic or tricyclic ring system having a N atom at the bridgehead and optionally containing 1, 2 or 3 additional heteroatoms selected from N, O or S;

R₂ is H, halogen, or a C₁-C₆alkyl, C₁-C₆alkoxy, C₃-C₇cycloalkyl, aryl or heteroaryl group each optionally substituted;

R₃ and R₄ are each independently H or an optionally substituted C₁-C₆alkyl group;

R₅ is H or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₇cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted;

R₆ is a C₁-C₆alkyl, C₃-C₇cycloalkyl, C₂-C₆alkenyl or C₂-C₆alkynyl group each optionally substituted;

R₇ and R₈ are each independently H or a C₁-C₆alkyl, C₃-C₇cycloalkyl, C₂-C₆alkenyl or C₂-C₆alkynyl group each optionally substituted;

m, n and p are each independently 0 or an integer of 1, 2 or 3;

q and x are each independently 0 or an integer of 1 or 2;

R₉, R₁₀, R₁₃ and R₁₇ are each independently H or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₆cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted;

R_{11} and R_{12} are each independently H or an optionally C_1 - C_6 alkyl group or R_{11} and R_{12} may be taken together with the atom to which they are attached to form a 5- to 7-member ring optionally containing another heteroatom selected from O, N or S;

R_{14} and R_{15} are each independently H or an optionally substituted C_1 - C_4 alkyl group or R_{14} and R_{15} may be taken together with the atom to which they are attached to form a 5- to 7-membered ring optionally containing another heteroatom selected from O, NR_{18} or SO_x ;

R_{16} is a C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_7 cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted; and

R_{18} is H or a C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_7 cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted; or

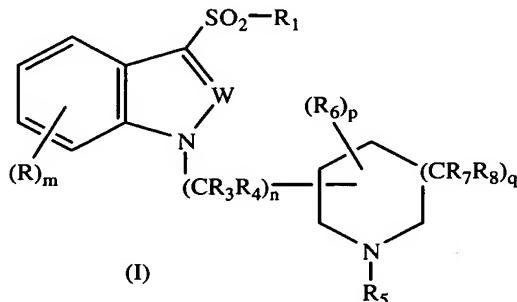
the stereoisomers thereof or the pharmaceutically acceptable salts thereof.

2. (Original) The compound according to claim 1 wherein n is 0.
3. (Original) The compound according to claim 1 wherein R_5 is H.
4. (Original) The compound according to claim 1 wherein R_1 is an optionally substituted phenyl group.
5. (Original) The compound according to claim 2 wherein q is 0 or 1.
6. (Original) The compound according to claim 2 wherein m is 0 and p is 0.
7. (Original) The compound according to claim 5 wherein the piperidinyl or pyrrolidinyl group is attached in the 3-position.
8. (Original) The compound according to claim 6 wherein R_1 is an optionally substituted phenyl group and q is 0 or 1.
9. (Cancelled)
10. (Cancelled)
11. (Cancelled)
12. (Cancelled)

13. (Cancelled)

14. (Cancelled)

15. (Currently Amended) A pharmaceutical composition which comprises a pharmaceutically acceptable carrier and an effective amount of a compound of formula I



wherein

W is [[N or]] CR₂;

R is halogen, CN, OCO₂R₉, CO₂R₁₀, CONR₁₁R₁₂, SO_xR₁₃, NR₁₄R₁₅, OR₁₆, COR₁₇ or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₇cycloalkyl, aryl or heteroaryl group each optionally substituted;

R₁ is an optionally substituted C₁-C₆alkyl, C₃-C₇cycloalkyl, aryl, or heteroaryl group or an optionally substituted 8- to 13-membered bicyclic or tricyclic ring system having a N atom at the bridgehead and optionally containing 1, 2 or 3 additional heteroatoms selected from N, O or S;

R₂ is H, halogen, or a C₁-C₆alkyl, C₁-C₆alkoxy, C₃-C₇cycloalkyl, aryl or heteroaryl group each optionally substituted;

R₃ and R₄ are each independently H or an optionally substituted C₁-C₆alkyl group;

R₅ is H or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₇cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted;

R₆ is a C₁-C₆alkyl, C₃-C₇cycloalkyl, C₂-C₆alkenyl or C₂-C₆alkynyl group each optionally substituted;

R₇ and R₈ are each independently H or a C₁-C₆alkyl, C₃-C₇cycloalkyl, C₂-C₆alkenyl or C₂-C₆alkynyl group each optionally substituted;

m, n and p are each independently 0 or an integer of 1, 2 or 3;

q and x are each independently 0 or an integer of 1 or 2;

R₉, R₁₀, R₁₃ and R₁₇ are each independently H or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, C₃-C₆cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted;

R_{11} and R_{12} are each independently H or an optionally C_1 - C_6 alkyl group or R_{11} and R_{12} may be taken together with the atom to which they are attached to form a 5- to 7-member ring optionally containing another heteroatom selected from O, N or S;

R_{14} and R_{15} are each independently H or an optionally substituted C_1 - C_4 alkyl group or R_{14} and R_{15} may be taken together with the atom to which they are attached to form a 5- to 7-membered ring optionally containing another heteroatom selected from O, NR_{18} or SO_x ;

R_{16} is a C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_7 cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted; and

R_{18} is H or a C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_7 cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted; or

the stereoisomers thereof or the pharmaceutically acceptable salts thereof.

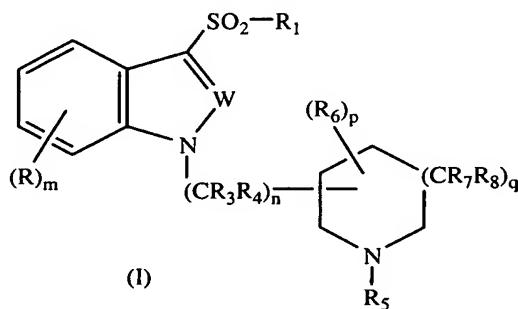
16. (Original) The composition according to claim 15 having a formula I compound wherein n is 0.

17. (Original) The composition according to claim 16 having a formula I compound wherein R_5 is H and q is 0 or 1.

18. (Original) The composition according to claim 17 having a formula I compound wherein R_1 is an optionally substituted phenyl group.

19. (Original) The composition according to claim 18 having a formula I compound wherein the piperidinyl or pyrrolidinyl group is attached in the 3-position.

20. (Currently Amended) A process for the preparation of a compound of formula I



wherein

W is [[N or]] CR_2 ;

R is halogen, CN, OCO_2R_9 , CO_2R_{10} , $\text{CONR}_{11}\text{R}_{12}$, SO_xR_{13} , $\text{NR}_{14}\text{R}_{15}$, OR_{16} , COR_{17} or a $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_2\text{-C}_6$ alkenyl, $\text{C}_2\text{-C}_6$ alkynyl, $\text{C}_3\text{-C}_7$ cycloalkyl, aryl or heteroaryl group each optionally substituted;

R_1 is an optionally substituted $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_3\text{-C}_7$ cycloalkyl, aryl, or heteroaryl group or an optionally substituted 8- to 13-membered bicyclic or tricyclic ring system having a N atom at the bridgehead and optionally containing 1, 2 or 3 additional heteroatoms selected from N, O or S;

R_2 is H, halogen, or a $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_6$ alkoxy, $\text{C}_3\text{-C}_7$ cycloalkyl, aryl or heteroaryl group each optionally substituted;

R_3 and R_4 are each independently H or an optionally substituted $\text{C}_1\text{-C}_6$ alkyl group;

R_5 is H or a $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_2\text{-C}_6$ alkenyl, $\text{C}_2\text{-C}_6$ alkynyl, $\text{C}_3\text{-C}_7$ cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted;

R_6 is a $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_3\text{-C}_7$ cycloalkyl, $\text{C}_2\text{-C}_6$ alkenyl or $\text{C}_2\text{-C}_6$ alkynyl group each optionally substituted;

R_7 and R_8 are each independently H or a $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_3\text{-C}_7$ cycloalkyl, $\text{C}_2\text{-C}_6$ alkenyl or $\text{C}_2\text{-C}_6$ alkynyl group each optionally substituted;

m , n and p are each independently 0 or an integer of 1, 2 or 3;

q and x are each independently 0 or an integer of 1 or 2;

R_9 , R_{10} , R_{13} and R_{17} are each independently H or a $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_2\text{-C}_6$ alkenyl, $\text{C}_2\text{-C}_6$ alkynyl, $\text{C}_3\text{-C}_6$ cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted;

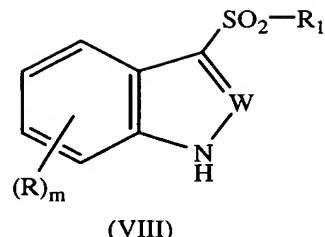
R_{11} and R_{12} are each independently H or an optionally $\text{C}_1\text{-C}_6$ alkyl group or R_{11} and R_{12} may be taken together with the atom to which they are attached to form a 5- to 7-membered ring optionally containing another heteroatom selected from O, N or S;

R_{14} and R_{15} are each independently H or an optionally substituted $\text{C}_1\text{-C}_4$ alkyl group or R_{14} and R_{15} may be taken together with the atom to which they are attached to form a 5- to 7-membered ring optionally containing another heteroatom selected from O, NR_{18} or SO_x ;

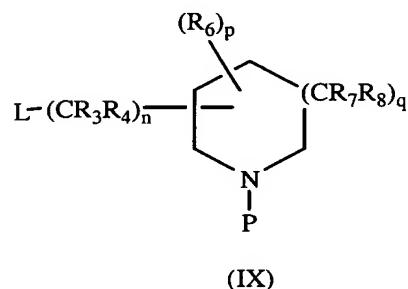
R_{16} is a $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_2\text{-C}_6$ alkenyl, $\text{C}_2\text{-C}_6$ alkynyl, $\text{C}_3\text{-C}_7$ cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted; and

R_{18} is H or a $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_2\text{-C}_6$ alkenyl, $\text{C}_2\text{-C}_6$ alkynyl, $\text{C}_3\text{-C}_7$ cycloalkyl, cycloheteroalkyl, aryl or heteroaryl group each optionally substituted

which process comprises reacting a compound of formula VIII



wherein W, R, R₁ and m are as described hereinabove with a protected azacyclic compound of formula IX



wherein P is a protecting group; L is a leaving group; and R₃, R₄, R₆, R₇, R₈, n, p and q are as described hereinabove in the presence of a first base to give the protected formula I compound; and deprotecting said compound to give the free amine of formula I wherein R₅ is H optionally alkylating said amine with an alkylating agent, R₅-L', wherein L' is a leaving group in the presence of a second base.